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| **PI NAME** |  |
| **STUDY TITLE** |  |

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| **1.** | What are you requesting? (***Choose ONE***) | |
| ☐ | Waiver of the requirement to obtain informed consent [45 CFR 46.116(f)(1)] |
| ☐ | Alteration of one or more of the required elements of informed consent. [45 CFR 46.116(f)(2)] |
|  | ☐ | Waiver of documentation of informed consent |
| If requesting a Waiver of Documentation of Consent, why? | | |
|  | ☐ | The informed consent form would be the principal risk and potentially harm participants through a breach of confidentiality. |
|  | ☐ | The research adds no more than minimal risk of harm to subjects, and written consent is normally not expected outside the research context. |
|  | ☐ | Participants are part of a group or community where signing forms is not the norm, and the research adds no more than minimal risk of harm to subjects. |
| If requesting a Waiver of Documentation of Consent, how will you get consent? | | |
|  | ☐ | The subject will provide consent verbally and be given an information sheet. (Submit Information Sheet alongside IRB packet) |
|  | ☐ | A questionnaire/survey will be given to the subject, and the return of the questionnaire/survey will indicate consent. (Submit Information Sheet alongside IRB packet). |
|  | ☐ | The subject will be told verbally about the study and will provide consent verbally. (Submit a Consenting Script alongside IRB packet). |
|  | ☐ | The subject will read the consent via the Internet and indicate consent by selecting an “I agree” button or similar. (Submit Internet Consent Text for IRB review). |
|  | ☐ | Other: |
|  | | |
| **2.** | Is this request for (***Choose ONE***): | |
| ☐ | All subjects |
| ☐ | Some subjects. Identify the group of subjects and provide rationale: |
| Rationale for Request | | |
| **3.** |  | |

**Indicate how consent will be obtained (choose ALL that apply):**

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| **4.** | ☐ | The subject will be given an Information Sheet to read and will provide consent verbally. (Submit Information Sheet for IRB review.) |
| ☐ | An Information Sheet will be mailed with a questionnaire/survey to the subject, and the return of the questionnaire/survey will indicate consent. (Submit Information Sheet for IRB review.) |
| ☐ | The subject will be told verbally about the study and will provide consent verbally. (Submit a Consenting Script for IRB review.) |
| ☐ | The subject will read the consent via the Internet and indicate consent by selecting an “I agree” button or similar. (Submit Internet Consent Text for IRB review). |
| ☐ | Other. Please describe: |